



## Project Managers

KLIFO is expanding and wants to engage Project Managers into a dynamic and experienced team within Clinical Trial Services.

The people we want to engage like to work in a consulting environment and have a positive, proactive, flexible, self-driven and self-confident personality.

We can offer a highly flexible, free and trustful working climate with exciting projects among competent colleagues where your contribution is valuable and makes a difference.

KLIFO is a service provider of drug development competences and operational services to the pharmaceutical, biotech and med-tech companies in Europe, US and Asia. Our services include Clinical Trial Supply, Clinical Trial Services, Regulatory Affairs, Pharmacovigilance, Chemistry and Manufacturing Control and Drug Development Counselling.

### The position as Project Manager:

The Project Manager is overall responsible for managing the clinical trials, i.e.:

- Management of assigned project in all phases, from start-up to the end of the trial
- Leadership of people involved in the project
- Investigational site selection in order to assure the quality and conformity of the sites
- Attending cross-functional project team in relation to trial(s)
- Operational and scientific input to key project documents
- Continuous relationship with the Principal Investigators and Sponsor to assure the success of the trial in terms of enrolment and quality
- Management and resolution of issues occurring during the trial
- Generation and management of Project Plans
- Investigators and Monitors training and organization of meetings
- Elaboration of trial specific procedures
- Participation in data management activities

### Education, experiences, knowledge and skill:

The Project Manager should possess the following qualifications:

- MSc in the life sciences field and a minimum of 5-6 years of overall experience out of which 2 years must be in project management in the pharmaceutical industry /Biotech/CRO
- Extensive knowledge of GCP guidelines, applicable regulatory requirements, medical terminology and clinical trial processes
- Willingness to travel
- Demonstrated project management skills including the ability to plan a project and operate within plan and budget
- Excellent verbal and written communication skills
- Computer skills, ability to develop and maintain excel spreadsheets and to generate PowerPoint presentations

In addition to the above-mentioned qualifications, the candidate is a dedicated and collaborative team player that possesses excellent planning skills and is fluent, spoken and written, in English and in a Scandinavian language.

### We offer:

- Work within different therapeutic areas and with tasks of varying complexity
- Work with a heterogeneous client pool
- Build international client relations
- Use and elaborate your competences and experience
- Work in an interactive, flexible and positive working environment

As KLIFO is a smaller service provider you will experience a higher level of transparency, influence and a good possibility for individual planning.

### Location:

KLIFO is located at Smedeland 36, 2600 Glostrup.

### Contact:

For more information, please contact Klas Rådberg, Clinical Research Director of CTS at +45 44 222 935

### Applications should be sent to:

[Mette.Widen@klifo.com](mailto:Mette.Widen@klifo.com), marked Project Manager

### Deadline:

24th August, 2018

For more information on KLIFO, please visit: [www.klifo.com](http://www.klifo.com)